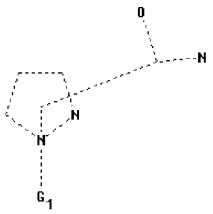
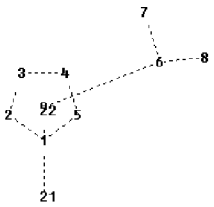
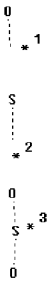
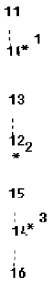
=>
Uploading C:\Program Files\Stnexp\Queries\10547996-broad.str









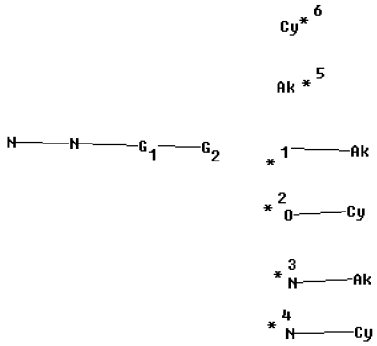
chain nodes :
6 7 10 11 12 13 14 15 16 21
ring nodes :
1 2 3 4 5
ring/chain nodes :
8
chain bonds :
6-7 6-8 10-11 12-13 14-15 14-16
ring bonds :
1-2 1-5 2-3 3-4 4-5
exact/norm bonds :
1-2 1-5 2-3 3-4 4-5 6-7 6-8 10-11 12-13 14-15 14-16
isolated ring systems :
containing 1 :

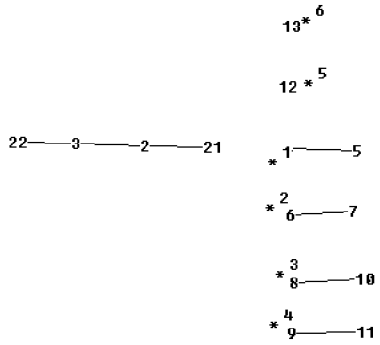
G1:[*1],[*2],[*3]

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS
11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 21:CLASS 22:CLASS

L1 STRUCTURE UPLOADED

=>
Uploading C:\Program Files\Stnexp\Queries\10547996-narrow-2.str





chain nodes :
2 4 5 6 7 8 9 10 11 12 13 21
ring nodes :
3 22
chain bonds :
2-3 2-21 4-5 6-7 8-10 9-11
ring bonds :
3-22
exact/norm bonds :
2-3 2-21 3-22 4-5 6-7 8-10 9-11

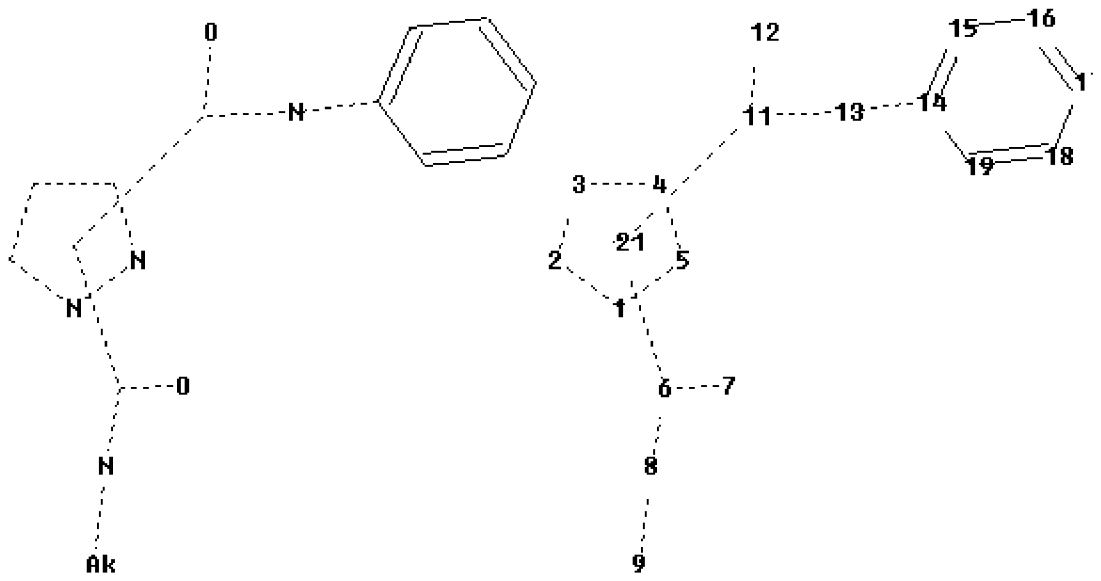
G1:C,S

G2:[*1],[*2],[*3],[*4],[*5],[*6]

Match level :
2:CLASS 3:Atom 4:CLASS 5:CLASS 6:CLASS 7:Atom 8:CLASS 9:CLASS 10:CLASS
11:Atom
12:CLASS 13:Atom 21:CLASS 22:Atom

L6 STRUCTURE UPLOADED

=>
Uploading C:\Program Files\Stnexp\Queries\10547996-elected.str



chain nodes :
6 7 8 9 11 12 13
ring nodes :
1 2 3 4 5 14 15 16 17 18 19
chain bonds :
6-7 6-8 8-9 11-12 11-13 13-14
ring bonds :
1-2 1-5 2-3 3-4 4-5 14-15 14-19 15-16 16-17 17-18 18-19
exact/norm bonds :
1-2 1-5 2-3 3-4 4-5 6-7 6-8 8-9 11-12 11-13 13-14
normalized bonds :
14-15 14-19 15-16 16-17 17-18 18-19
isolated ring systems :
containing 14 :

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:Atom
11:CLASS 12:CLASS 13:CLASS 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
21:Atom

L9 STRUCTURE UPLOADED

FILE 'REGISTRY' ENTERED AT 11:31:50 ON 03 JUN 2008
L1 STRUCTURE UPLOADED
L3 2653 S L1 SSS FULL

L6 STRUCTURE UPLOADED
L8 2616 S L6 SSS FULL SUB=L3
SAV TEM L8 NAR547996/A

L9 STRUCTURE UPLOADED
L11 46 S L9 SSS FULL SUB=L8

FILE 'CAPLUS' ENTERED AT 11:43:43 ON 03 JUN 2008
L12 4 S L11
L13 1 S US200!-547996/APPS

L14 1 S L12 AND L13
L15 3 S L12 NOT L13

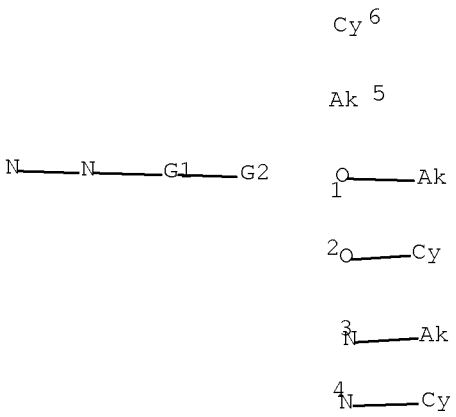
FILE 'REGISTRY' ENTERED AT 11:44:07 ON 03 JUN 2008

=> d l1
L1 HAS NO ANSWERS
L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

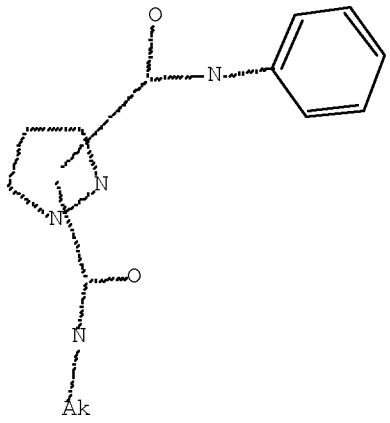
=> d l6
L6 HAS NO ANSWERS
L6 STR



G1 C,S
G2 [@1],[@2],[@3],[@4],[@5],[@6]

Structure attributes must be viewed using STN Express query preparation.

=> d l9
L9 HAS NO ANSWERS
L9 STR



Structure attributes must be viewed using STN Express query preparation.

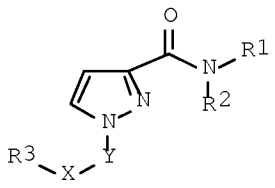
=> fil caplus

=> d l14 bib abs

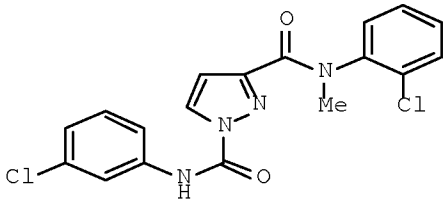
L14 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2004:780692 CAPLUS Full-text
DN 141:296011
TI Preparation of pyrazoles having 15-lipoxygenase inhibitory activity useful for treating inflammation
IN Hallberg, Anders; Schaal, Wesley; Larhed, Mats; Olofsson, Kristofer; Pelcman, Benjamin; Sanin, Andrei
PA Biolipox AB, Swed.; McNeeney, Stephen Phillip
SO PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004080999	A1	20040923	WO 2004-GB1054	20040312
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	EP 1603897	A1	20051214	EP 2004-720088	20040312
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK				
	JP 2006520373	T	20060907	JP 2006-505952	20040312
	US 20060183780	A1	20060817	US 2005-547996	20051123 <--
PRAI	SE 2003-705	A	20030314		

US 2003-482563P P 20030626
WO 2004-GB1054 W 20040312
OS MARPAT 141:296011
GI



I



II

AB Tilte compds. I [wherein R1 = (un)substituted hetero/aryl; R2 = H, alkyl optionally substituted by halo; or when R2 = alkyl optionally substituted by halo, R1NR2 = (un)substitytuted 5- to 7-membered ring, optionally containing 1 to 3 heteroatoms, and/or 1 to 3 double bonds; R3 = (un)substituted alk(en/yn)yl, hetero/cycloalkyl, hetero/aryl; X = a direct bond, O, NH and derivs.; Y = C(:O), C(:S), SO2; and their pharmaceutically-acceptable salts; with provisos excluding certain compds.] were prepared The use of compds. I (without exclusions) as inhibitors of the activity of a lipoxygenase, in particular 15-lipoxygenase, for treating inflammation is claimed. Thus, reacting 1H-Pyrazole-3-carboxylic acid N-(2-chlorophenyl)-N-methylamide (preparation given) with 3-chlorophenylisocyanate in toluene at 100° gave dicarboxamide II. Five representative compds. gave inhibition of 15-lipoxygenase ranging from 39 to 50% in a fluorescence bioassay.
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d l15 tot bib abs hitstr

L15 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2008:550983 CAPLUS Full-text
DN 148:517709
TI Preparation of substituted pyrazolecarboxanilide derivatives or salts thereof as agricultural or horticultural chemicals
IN Machiya, Kozo; Matsuzaki, Yoshihiro; Furuya, Takashi; Suwa, Akiyuki; Yasokawa, Noriaki; Fujioka, Shinsuke
PA Nihon Nohyaku Co., Ltd., Japan
SO PCT Int. Appl., 86pp.
CODEN: PIXXD2
DT Patent
LA Japanese

FAN.CNT 1

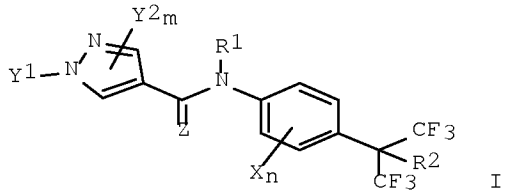
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2008053991	A1	20080508	WO 2007-JP71403	20071102

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRAI JP 2006-299561 A 20061102

GI



I

AB The title compds. [I; R1 = H, alkyl, alkylcarbonyl, alkenylcarbonyl, cycloalkyl, (substituted) phenylalkyl, (substituted) phenylcarbonyl, etc.; R2 = H, halogeno, alkyl, cyano, OH, alkoxy, (substituted) phenoxy, (substituted) phenylthio, (substituted) phenylsulfonyl, etc.; Z = O or S; X = H, halogeno, CN, alkyl, etc.; Y1 = alkylcarbonyl, cycloalkylcarbonyl, alkoxyalkyl, alkoxycarbonyl, (substituted) phenoxycarbonyl, alkoxyalkylcarbonyl, alkylcarbonyloxyalkyl, alkylsulfonyl, (substituted) phenylsulfonyl, mono- or dialkylaminocarbonyl, mono- or dialkylaminothiocarbonyl, (substituted) phenylcarbonyl, (substituted) phenylalkyl, dialkoxy(thio)phosphoryl, etc.; Y2 = halogeno, cyano, NO2, alkyl, (substituted) Ph, (substituted) phenoxy, etc.; m = 1 or 2; n = 1-4] or salts thereof are prepared These compds. are useful as agricultural or horticultural chems., in particular insecticides or acaricides. Thus, 0.06 g acetyl chloride was added to a suspension of 0.30 g N-isobutyryl-N-[3-isobutyl-4-[1-methoxy-2,2,2-trifluoro-1-(trifluoromethyl)ethyl]phenyl]-1H-3,5-dimethylpyrazole-4-carboxamide and 0.12 g K2CO3 in 10 mL MeCN and the resulting mixture was stirred at room temperature for 3 h to give 76.2% N-isobutyryl-N-[3-isobutyl-4-[1-methoxy-2,2,2- trifluoro-1-(trifluoromethyl)ethyl]phenyl]-1-acetyl-3,5-dimethylpyrazole-4- carboxamide (II). II at 500 ppm controlled 90-99% adult Tetranychus urticae.

IT 1022987-62-7P, N-Isobutyryl-N-[3-isobutyl-4-[1-methoxy-2,2,2-trifluoro-1-(trifluoromethyl)ethyl]phenyl]-1-(dimethylcarbamoyl)-3,5-dimethylpyrazole-4-carboxamide

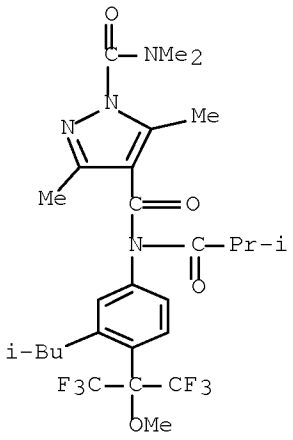
RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of substituted pyrazolecarboxanilide derivs. or salts thereof

as agricultural or horticultural chems. such as insecticides and acaricides)

RN 1022987-62-7 CAPLUS

CN INDEX NAME NOT YET ASSIGNED



RE.CNT 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L15 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2008:10602 CAPLUS Full-text

DN 148:113191

TI Methods for identifying modulators of Eoxin formation

IN Claesson, Hans-Erik; Bjoerkholm, Magnus

PA Biolipox AB, Swed.; Pilkington, Stephanie

SO PCT Int. Appl., 126pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2008001079	A1	20080103	WO 2007-GB2394	20070627

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

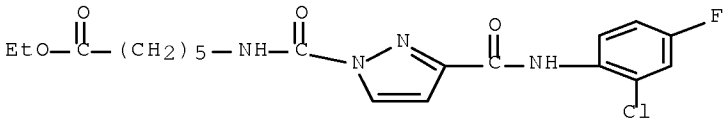
PRAI SE 2006-1394 A 20060627

AB A method for identifying a compound for modulating the formation of 14,15-LTC4 (Eoxin C4; EoxC4), 14,15-LTD4 (Eoxin D4; EoxD4), or 14,15-LTE4 (Eoxin E4; EoxE4) in a biol. system. A method for identifying a compound with an anti-inflammatory effect, the method comprising testing the compound for an effect on formation and/or activity of 14,15-LTC4 (Eoxin C4; EoxC4), 14,15-LTD4

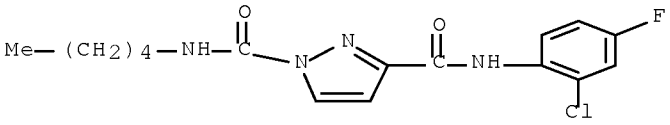
(Eoxin D4; EoxD4), or 14,15-LTE4 (Eoxin E4; EoxE4) in a biol. system. A method of making an anti-inflammatory composition or Eoxin formation-modulating composition comprising (i) identifying an anti-inflammatory compound or Eoxin formation-modulating compound by a method of the invention; (ii) combining the compound with a pharmaceutically acceptable excipient or carrier.

IT 763108-23-2 763108-24-3 1000678-87-4
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(Eoxin formation inhibition by; methods for identifying modulators of Eoxin formation as anti-inflammatory agents and bone loss inhibitors and for Eoxins to promote inflammation)

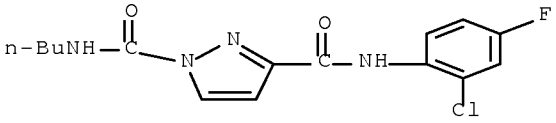
RN 763108-23-2 CAPLUS
CN Hexanoic acid, 6-[[[3-[[(2-chloro-4-fluorophenyl)amino]carbonyl]-1H-pyrazol-1-yl]carbonyl]amino]-, ethyl ester (CA INDEX NAME)



RN 763108-24-3 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-pentyl- (CA INDEX NAME)



RN 1000678-87-4 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N1-butyl-N3-(2-chloro-4-fluorophenyl)- (CA INDEX NAME)



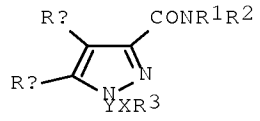
RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L15 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2006:295966 CAPLUS Full-text
DN 144:350669
TI Preparation of pyrazolecarboxamides as 15-lipoxygenase inhibitors for treatment of inflammation.

IN Pelcman, Benjamin; Sanin, Andrei; Nilsson, Peter; Boesen, Thomas
PA Biolipox AB, Swed.
SO PCT Int. Appl., 87 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006032852	A1	20060330	WO 2005-GB3584	20050919
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	EP 1794130	A1	20070613	EP 2005-784086	20050919
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
	JP 2008513427	T	20080501	JP 2007-531832	20050919
	US 20080090836	A1	20080417	US 2007-663180	20070319
PRAI	US 2004-610952P	P	20040920		
	WO 2005-GB3584	W	20050919		

OS MARPAT 144:350669
GI



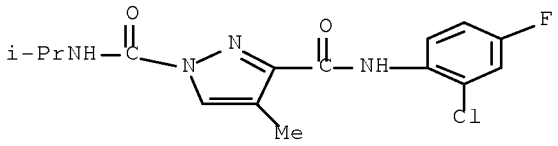
I

AB Title compds. [I; R1 = (substituted) aryl, heteroaryl; R2 = (substituted) alkyl; R3 = (substituted) alkyl, heterocycloalkyl, aryl, heteroaryl; X = bond, NR4a; Y = CO, CS, SO2; R4a = H, (substituted) alkyl, heterocycloalkyl; Ra, Rb = H, halo, (substituted) alkyl], were prepared Thus, 5-methylpyrazole-1,3-dicarboxylic acid 3-[(2-chloro-4- fluorophenyl)amide]-1-hexylamide (preparation outlined) inhibited 15-lipoxygenase with IC50 = 0.40 μM.

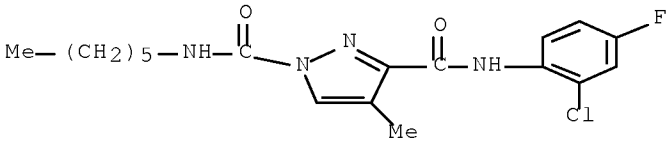
IT 881683-98-3P 881684-00-0P 881684-01-1P
881684-02-2P 881684-03-3P 881684-04-4P
881684-06-6P 881684-09-9P 881684-10-2P
881684-11-3P 881684-12-4P 881684-13-5P
881684-14-6P 881684-17-9P 881684-18-0P
881684-21-5P 881684-22-6P 881684-23-7P
881684-24-8P 881684-25-9P 881684-26-0P
881684-27-1P 881684-28-2P 881684-32-8P
881684-81-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(Uses)
(preparation of pyrazolecarboxamides as 15-lipoxygenase inhibitors)

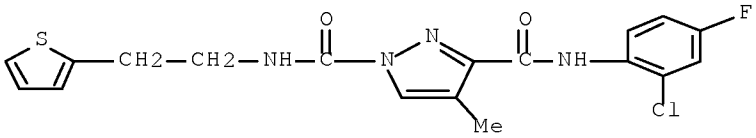
RN 881683-98-3 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-4-methyl-N1-(1-methylethyl)- (CA INDEX NAME)



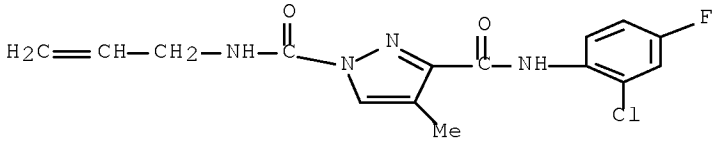
RN 881684-00-0 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-hexyl-4-methyl- (CA INDEX NAME)



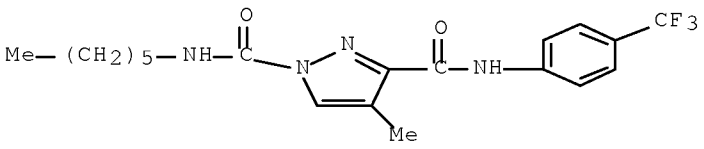
RN 881684-01-1 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-4-methyl-N1-[2-(2-thienyl)ethyl]- (CA INDEX NAME)



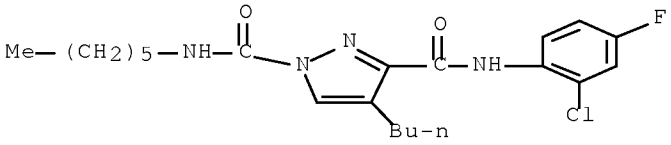
RN 881684-02-2 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-4-methyl-N1-2-propen-1-yl- (CA INDEX NAME)



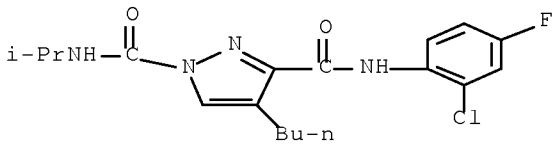
RN 881684-03-3 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N1-hexyl-4-methyl-N3-[4-(trifluoromethyl)phenyl]- (CA INDEX NAME)



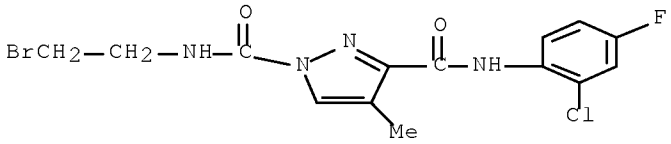
RN 881684-04-4 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, 4-butyl-N3-(2-chloro-4-fluorophenyl)-N1-hexyl- (CA INDEX NAME)



RN 881684-06-6 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, 4-butyl-N3-(2-chloro-4-fluorophenyl)-N1-(1-methylethyl)- (CA INDEX NAME)

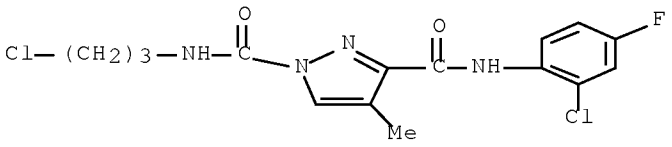


RN 881684-09-9 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N1-(2-bromoethyl)-N3-(2-chloro-4-fluorophenyl)-4-methyl- (CA INDEX NAME)

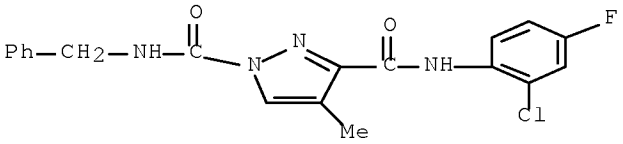


RN 881684-10-2 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-(3-

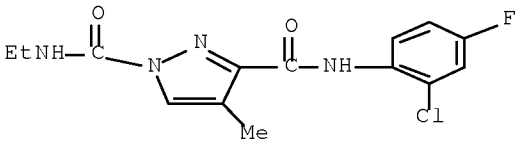
chloropropyl)-4-methyl- (CA INDEX NAME)



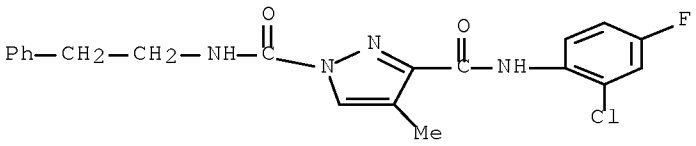
RN 881684-11-3 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-4-methyl-N1-(phenylmethyl)- (CA INDEX NAME)



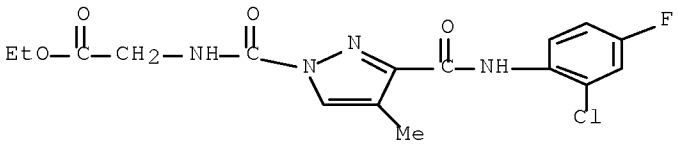
RN 881684-12-4 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-ethyl-4-methyl- (CA INDEX NAME)



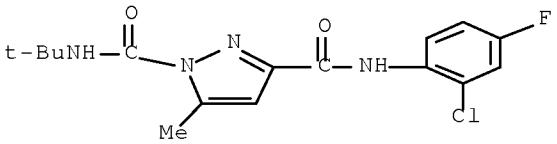
RN 881684-13-5 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-4-methyl-N1-(2-phenylethyl)- (CA INDEX NAME)



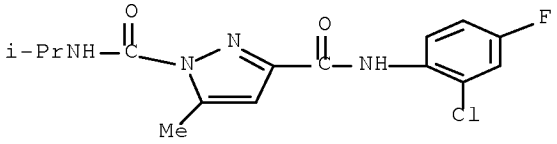
RN 881684-14-6 CAPLUS
CN Glycine, N-[[3-[[(2-chloro-4-fluorophenyl)amino]carbonyl]-4-methyl-1H-pyrazol-1-yl]carbonyl]-, ethyl ester (CA INDEX NAME)



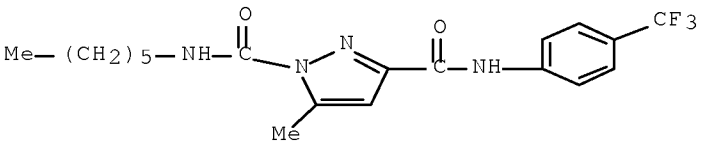
RN 881684-17-9 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-(1,1-dimethylethyl)-5-methyl- (CA INDEX NAME)



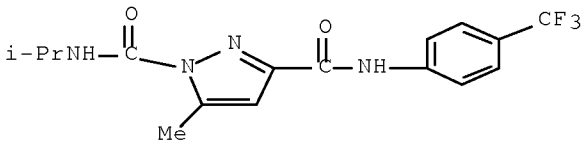
RN 881684-18-0 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-5-methyl-N1-(1-methylethyl)- (CA INDEX NAME)



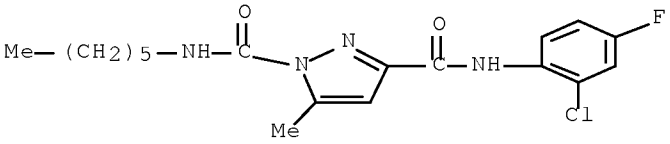
RN 881684-21-5 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N1-hexyl-5-methyl-N3-[4-(trifluoromethyl)phenyl]- (CA INDEX NAME)



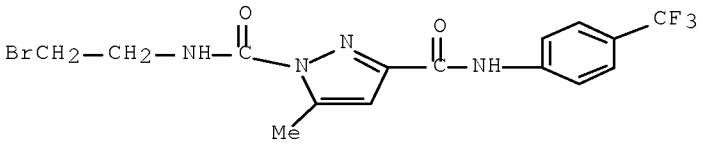
RN 881684-22-6 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, 5-methyl-N1-(1-methylethyl)-N3-[4-(trifluoromethyl)phenyl]- (CA INDEX NAME)



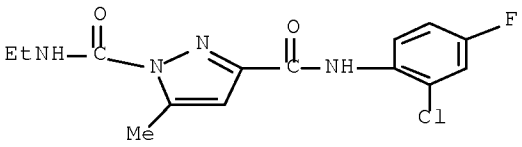
RN 881684-23-7 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-hexyl-5-methyl- (CA INDEX NAME)



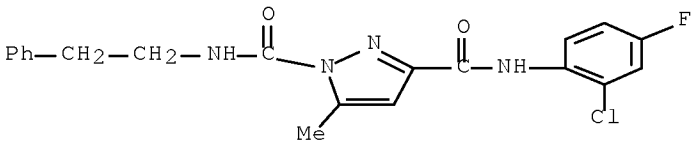
RN 881684-24-8 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N1-(2-bromoethyl)-5-methyl-N3-[4-(trifluoromethyl)phenyl]- (CA INDEX NAME)



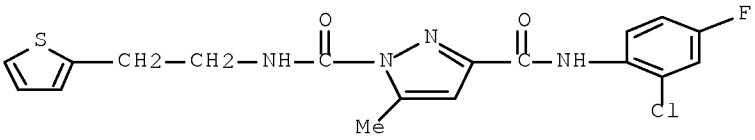
RN 881684-25-9 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-ethyl-5-methyl- (CA INDEX NAME)



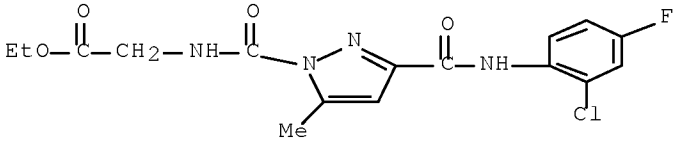
RN 881684-26-0 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-5-methyl-N1-(2-phenylethyl)- (CA INDEX NAME)



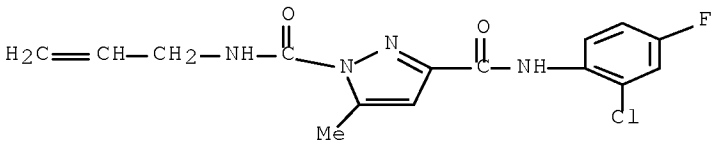
RN 881684-27-1 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-5-methyl-N1-[2-(2-thienyl)ethyl]- (CA INDEX NAME)



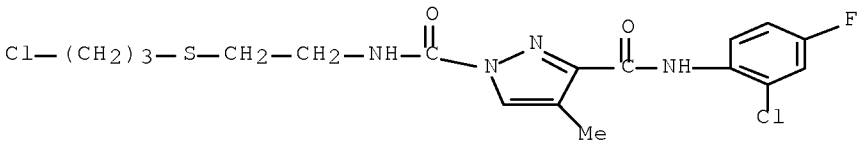
RN 881684-28-2 CAPLUS
CN Glycine, N-[[3-[[(2-chloro-4-fluorophenyl)amino]carbonyl]-5-methyl-1H-pyrazol-1-yl]carbonyl]-, ethyl ester (CA INDEX NAME)



RN 881684-32-8 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-5-methyl-N1-2-propen-1-yl- (CA INDEX NAME)



RN 881684-81-7 CAPLUS
CN 1H-Pyrazole-1,3-dicarboxamide, N3-(2-chloro-4-fluorophenyl)-N1-[2-[(3-chloropropyl)thio]ethyl]-4-methyl- (CA INDEX NAME)



RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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